ABSTRACT

Method of use of (imidazol-5-yl)methyl-2-quinolinone derivatives to inhibit smooth muscle cell proliferation.

This invention comprises the use of compounds of formula (I)

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wherein the dotted line represents an optional bond; X is oxygen or sulfur; R¹ is hydrogen, C₁₋₁₂alkyl, Ar¹, Ar²C₁₋₆alkyl, quinolinylC₁₋₆alkyl, pyridylC₁₋₆alkyl, hydroxyC₁₋₆alkyl, C₁₋₆alkyloxyC₁₋₆alkyl, mono- or di(C₁₋₆alkyl)aminoC₁₋₆alkyl, aminoC₁₋₆alkyl, or a radical of formula -Alk¹-C(=O)-R⁹, -Alk¹-S(O)-R⁹ or - $Alk^1-S(O)_2-R^9$; R^2 , R^3 and R^{16} each independently are hydrogen, hydroxy, halo, 15 cyano, C_{1-6} alkyloxy, hydroxy C_{1-6} alkyloxy, C_{1-6} alkyloxy C_{1-6} alkyloxy, amino C_{1-6} alkyloxy, mono- or di(C_{1-6} alkyl)amino C_{1-6} alkyloxy, Ar¹, Ar² C_{1-6} alkyl, Ar^2oxy , $Ar^2C_{1-6}alkyloxy$, hydroxycarbonyl, $C_{1-6}alkyloxycarbonyl$, trihalomethyl, trihalomethoxy, C₂₋₆alkenyl; R⁴ and R⁵ each independently are hydrogen, halo, Ar¹, 20 C_{1-6} alkyl, hydroxy C_{1-6} alkyl, C_{1-6} alkyloxy C_{1-6} alkyl, C_{1-6} alkyloxy, C_{1-6} alkylthio, amino, hydroxycarbonyl, C₁₋₆alkyloxycarbonyl, C₁₋₆alkylS(O)C₁₋₆alkyl or C₁₋₆alkylS(O)₂C₁₋₆alkyl; R⁶ and R⁷ each independently are hydrogen, halo, cyano, C_{1-6} alkyl, 4,4-dimethyl-oxazolyl, C_{1-6} alkyloxy or Ar²oxy; R⁸ is hydrogen, C_{1-6} alkyl, cyano, hydroxycarbonyl, C₁₋₆alkyloxycarbonyl, C₁₋₆alkylcarbonylC₁₋₆alkyl, cyanoC₁₋₆alkyl, C₁₋₆alkyloxycarbonylC₁₋₆alkyl, carboxyC₁₋₆alkyl, hydroxyC₁₋₆alkyl, 25 aminoC₁₋₆alkyl, mono- or di(C₁₋₆alkyl)aminoC₁₋₆alkyl, imidazolyl, haloC₁₋₆alkyl, C₁₋₆alkyloxyC₁₋₆alkyl, aminocarbonylC₁₋₆alkyl, or a radical of formula -O-R¹⁰, -S-R¹⁰, -N-R¹¹R¹²; R¹⁷ is hydrogen, halo, cyano, C₁₋₆alkyl, C₁₋₆alkyloxycarbonyl, Ar¹; R¹⁸ is hydrogen, C₁₋₆alkyl, C₁₋₆alkyloxy or halo; R¹⁹ is hydrogen or C₁₋₆alkyl; for the

manufacture of a medicament to inhibit smooth muscle cell proliferation.